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CLAIMS

- 1. A method of improving anesthesia recovery comprising the step of administering to an animal in need of such treatment a therapeutically effective amount of a pharmaceutical composition of a NK-1 receptor antagonist; a pharmaceutically acceptable salt thereof, a prodrug of said compound or said salt, or a solvate or hydrate of said compound, said salt or said prodrug.
- The use of a NK-1 receptor antagonist; a pharmaceutically acceptable salt
 thereof, a prodrug of said compound or said salt, or a solvate or hydrate of said compound, said salt or said prodrug, in the manufacture of a medicament for improving anesthesia.
 - 3. A method or use according to Claim 1 or Claim 2 wherein the NK-1 receptor antagonist is a compound of Formula I

wherein R^2 is selected from the group consisting of methyl, ethyl, isopropyl, *sec*-butyl and *tert*-butyl, or a pharmaceutically acceptable salt thereof.

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4. A method or use according to Claim 3 wherein the compound of Formula I is a compound of Formula Ia,

- 5 (2S,3S)-2-benzhydryl-*N*-(5-*tert*-butyl-2-methoxybenzyl)quinuclidin-3-amine, or a pharmaceutically acceptable salt thereof.
 - 5. The method or use according to Claim 4 wherein the compound is the citrate salt of the compound of Formula Ia.
 - 6. The method or use according to any previous claim wherein the composition is parenterally, enterally or orally administered prior, during or after an administration of a general anesthesia.
 - 7. The method or use according to Claim 6 wherein the composition is administered parenterally.
 - 8. The method or use according to Claim 7 wherein the composition further comprises a pharmaceutically acceptable cyclodextrin.
 - 9. The method or use according to Claim 7 or Claim 8 wherein the amount of the NK-1 antagonist is 0.01 mg/kg to 100 mg/kg of a patient's body weight.
- 10. A pharmaceutical composition for improving anesthesia recovery comprising a NK-1 receptor antagonist; a pharmaceutically acceptable salt thereof, a prodrug of said compound or said salt, or a solvate or hydrate of said compound, said salt or said prodrug.